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Plasmid DNA Isolation

Phage DNA

GIAGEN Instruments

Accessories

Appendices

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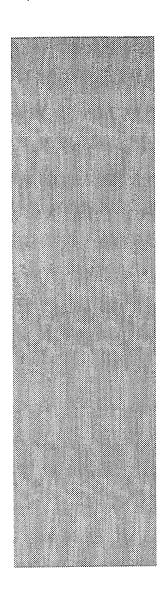
**QIAGEN Services** 

DNA Cleanup, Gel Extraction, and Bys-Terminator Removal

### WHAT'S NEW | PRODUCT GUIDE | PRODUCT LIST | HOW TO ORDER | TECHNICAL RESOURCES | TRADEMARKS & DISCLAIMERS | TALK TO GIAGEN | RNA Isolation and Purification giagen Click here for printer-friendly version Reverse Transcription and PCR RNeasy® Total RNA System Transfection The RNeasy® Total RNA isolation system is designed for fast and efficient isolation of Protein Expression. from a wide variety of samples. Different kits are tailored to suit different RNA purificati Purification. Detection, & Assay RNeasy sample preparation technology is fully licensed, allowing RNeasy purified nuc to be used in any molecular assay or other downstream application without risk of pate ANA isolation and Purification infringement. DMA toolation from Animals and Plants Features and benefits **DNA** and **RNA** tsolation from Clinical Samples

- High-quality total RNA in minutes
- No phenol/chloroform extraction, no CsCl gradients, no LiCl or ethanol precipita
- Excellent recovery of RNA from very small to large amounts of starting material
- Ready-to-use RNA for any downstream application

RNeasy Kit Options	and the second of the second o
Sample Source	Recommended RNeasy Kit
Animal cells Small-to large-scale 96-well format Automated, 96-well forn	RNeasy Mini, Midi, and Maxi Kits RNeasy 96 Kit natRNeasy 96 BioRobot Kit
Animal tissues Small- to large-scale	RNeasy Mini, Midi, and Maxi Kits
Bacteria or yeast Small-to large-scale	RNeasy Mini, Midi, and Maxi Kits
Whole blood	<u>see QiAamp<sup>®</sup> RNA Blood Mini Kit</u>
Plant cells or tissues	RNeasy Plant Mini Kit
Filamentous fungi	RNeasy Plant Mini Kit
In vitro transcripts and enzymatic reactions Small-to large-scale 96-well format	RNeasy Mini, Midi, and Maxi Kits RNeasy 96 Kit Additional RNeasy Buffers



### Principle ·

RNeasy Kits simplify total RNA isolation by combining the stringency of guanidine—isothiocyanate lysis with the speed of silica-gel—membrane purification. RNeasy technology replaces cumbersome and problematic RNA isolation procedures such as phenol/chloroform extraction\*, centrifugation through CsCl cushions, and precipitation with LiCl or alcohol. RNeasy purification also provides an enrichment for mRNA since most RNAs <200 nucleotides (which comprise 15–20% of total RNA) are selectively excluded.

#### Procedure ·

Samples are first lysed and then homogenized. Ethanol is added to the lysate to provide ideal binding conditions (see flowchart). The lysate is then loaded onto the RNeasy spin column. RNA binds, and all contaminants are efficiently washed away. Pure, concentrated RNA is eluted in water.

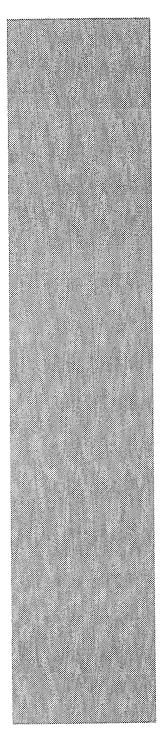
### Applications -

RNA purified with RNeasy technology has  $A_{260}/A_{280}$  ratios of 1.9–2.1<sup>†</sup> and is ideal for use in all applications. (See RNeasy references.) Downstream applications include:

- Northern, dot, and slot blotting
- RT-PCR
- Poly A<sup>+</sup> RNA selection



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#### Other sections of interest

References for total RNA isolation and cleanup with the RNeasy System

## Related Products

Total RNA isolation from whole blood with the QIAamp RNA Blood Mini Kit Viral RNA isolation from cell-free body fluids with the QIAamp Viral RNA Mini Kit

Poly A<sup>±</sup> mRNA isolation with the Oligotex™ System

Parallel isolation of RNA and genomic DNA and isolation of low-molecular-weight R

the QIAGEN® RNA/DNA System QIAGEN Reverse Transcriptases

#### Accessories

Homogenization of cell and tissue lysates with QlAshredder™ homogenizers DNase digestion during RNA purification with the RNase-Free DNase Set Plasticware and tape sheets

#### Technical information

Spectrophotometric quantitation of nucleic acids
Sizes and molecular weights of various RNAs
RNA content and distribution in various cells and tissues

QIAGEN purification technologies

#### **QIAGEN literature**

Brochure: High-Performance RNA

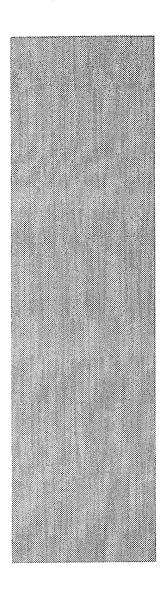
The QIAGEN RNA Club

\* RNA prepared using phenol-extraction methods may contain contaminating phenol, which affects the A<sub>260</sub> the purified RNA due to absorbance of phenol at 270 and 275 nm (see Stulnig, T.M. and Amberger, A. (1994 contaminating phenol in nucleic acid preparations. BioTechniques, **16**, 403).

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<sup>†</sup> Measured in 10 mM Tris-Cl, pH 7.5. See "Spectrophotometric Quantitation of Nucleic Acids."



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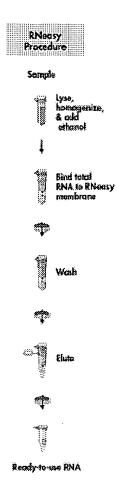
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# RNA Isolation and Purification

Reverse Transcription and PCR

Transfection

Protein Expression, Purification, Octoption, & Assay

ANA isolation and Purification

DMA Isolation from Animals and Plants

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# MINA ISOIATION AND LUMINOURIO

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# References for Total RNA Isolation and Cleanup with the RNeasy System

This is a partial list of papers citing RNeasy Kits for total RNA isolation and cleanup. Please contact QIAGEN Technical Services or your local distributor for a complete references.

### Selected RNeasy References

Dechant, G., Tsoulfas, P., Parada, L.F., and Barde, Y.A. (1997) The neurotrophin receptor p75 binds neurotrophin-3 on sympathetic neurons with high affinity and specificity. J. Neurosci. 17, 5281.

> Total RNA from chicken ganglia and neuronal cell cultures for RT-PCR

Dudareva, N. Cseka, L., Blanc, V.M., and Pichersky, E. (1996) Evolution of floral sc Clarkia: novel patterns of S-linalool synthase gene expression in the C. breweri flow Plant Cell 8, 1137.

Total RNA from Clarkia flower parts to localize gene expression

Estacio, W., Santa Anna-Arriola, S., Adedipe, M., and Márquez-Magaña, L.M. (1998 Dual promoters are responsible for transcription initiation of the flaiche operon in Ba subtilis. J. Bacteriol. 180, 3548.

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Gonzalez, P., Zigler, J.S., Jr., Epstein, D.L., and Borrás, T. (1999) Identific and isolation of differentially expressed genes from very small tissue sampl BioTechniques 26, 884.

▶ Total RNA from small amounts of human eye tissue for expression-arra analysis

Hassink, S.G. et al. (1997) Placental leptin: an important new growth facto intrauterine and neonatal development? Pediatrics 100(1), e1.

▶ Total RNA from human placenta and fat biopsy samples for multiplex RT-PCR

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Motlik, J., Carnwath, J.W., Herrmann, D. Terletski, V., Anger, M., and Niemann, H. (1998) Automated recording of RNA differential display patte from pig granulosa cells. BioTechniques 24, 148.

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Nakayama, J.-i., Tahara, H., Tahara, E., Saito, M., Ito, K., Nakamura, H., Nakanishi, T., Tahara, E., Ide, T., and Ishikawa, F. (1998) Telomerase activation by hTRT in human normal fibroblasts and hepatocellular carcino Nature Genet. 18, 65.

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Nichols, B.L., et al. (1997) Effects of malmitrition on expression and activit lactase in children. Gastroenterology 112, 742.

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Outinen, P.A., et al. (1998) Characterization of the stress-inducing effects o homocysteine. Biochem. J. 332, 213.

 RNeasy total RNA and Oligotex mRNA for expression-array and differential-display analysis

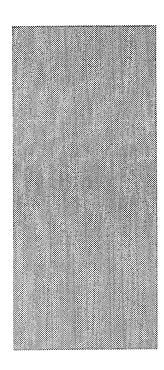
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Rieder, G., Hatz, R.A., Moran, A.P., Walz, A., Stolte, M., and Enders, G. (1 Role of adherence in interleukin-8 induction in Helicobacter pylori-associat



gastritis Infect. Immun. 65, 3622.

Total RNA from human gastric biopsy samples for competitive RT-PCR

Su, S., Vivier, R.G., et al. (1997) High-throughput RT-PCR analysis of mul transcripts using a microplate RNA isolation procedure. BioTechniques 22, 1107.

▶ High-throughput RNA isolation with the RNeasy 96 Kit for transcription-pattern analysis

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